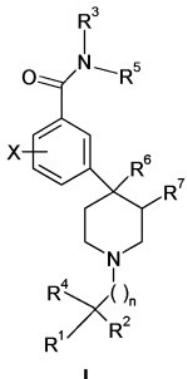


Claims:

1. (Currently Amended) A compound according to formula I:



wherein X is H, halogen, or CN;

R¹ and R² are independently H, C₁-C₆ alkyl, -(CH₂)_k-aryl, -(CH₂)_k-heteroaryl, wherein said alkyl, -(CH₂)_k-aryl or -(CH₂)_k-heteroaryl group is optionally substituted anywhere on said group with one or more R¹² groups, or, with the carbon to which R⁴ and R² are attached, are connected to form a C₃-C₇ cycloalkyl or a 4-7 membered carbocyclic or heterocycloalkyl comprising from one to three hetero moieties selected from O, S, C(=O), and N; and wherein said cycloalkyl or heterocycloalkyl optionally contains one or more double bonds; and wherein said cycloalkyl or heterocycloalkyl is optionally fused to or substituted with a C₆-C₁₄ aryl or 5-14 membered heteroaryl group; wherein said C₃-C₇ cycloalkyl or 4-7 membered carbocyclic or heterocycloalkyl formed by R⁴ and R² can each optionally be substituted by from one to three R¹² groups, and said optionally fused or substituted aryl or heteroaryl, substituted alkyl, substituted aryl optionally fused aryl or heteroaryl may each optionally independently be substituted with from one to six R¹² groups in any stereochemical relationship;

wherein the R¹² groups are independently selected from H, R¹³, R¹⁶, -C₁-C₄ alkyl optionally containing one or two unsaturated bonds, halogen, -OR¹³, -NO₂, -CN, -C₃-C₆ cycloalkyl, aryl, substituted aryl, wherein said aryl or substituted aryl is independently optionally substituted with 1-3 R¹⁸ groups, -C(R⁴)(C₁-C₄ alkyl)(C₁-C₄ alkyl) wherein said

alkyl groups may form a C₃-C₇ carbocyclic ring, -(CH₂)_v-NR¹³R¹⁴, -NR¹³C(=O)R¹⁴, -C(=O)NR¹³R¹⁴, -OC(=O)R¹³, -C(=O)OR¹³, -C(=O)R¹³, -NR¹³C(=O)OR¹⁴, -NR¹³C(=O)NR¹⁴R¹⁵, -NR¹³S(=O)R¹⁴, -NR¹⁷S(=O)NR¹³R¹⁴ and -S(=O)R¹³;

R¹⁸ is H, F, Cl, -OH, -C₁-C₄ alkyl, -C≡N, -NR¹³C(=O)R¹⁴, -C(=O)NR¹³R¹⁴, -O(C₁-C₄)alkyl, -NH(C₁-C₄)alkyl, -N(C₁-C₄)alkyl(C₁-C₄)alkyl, -(CH₂)_nOH, -(CH₂)_nC≡N, -(CH₂)_n-NR¹³C(=O)R¹⁴, -(CH₂)_n-C(=O)NR¹³R¹⁴, -(CH₂)_n-O(C₁-C₄)alkyl, -(CH₂)_n-NH₂, -(CH₂)_n-NH(C₁-C₄)alkyl or -(CH₂)_n-N(C₁-C₄)alkyl(C₁-C₄)alkyl;

R⁴ is absent or is H, -C₁-C₄ alkyl which may optionally contain one or two unsaturated bonds, -OH, O-(C₁-C₄)alkyl, (C₁-C₄)-alkyl-OH, (CH₂)_nNH₂, -(CH₂)_n-NH(C₁-C₄)alkyl, (CH₂)_n-N(C₁-C₄)alkyl(C₁-C₄)alkyl, -(CH₂)_n-NHC(=O)(C₁-C₄)alkyl, -(CH₂)_n-NO₂, -(CH₂)_n-C≡N, -(CH₂)_n-C(=O)NH₂, -(CH₂)_n-C(=O)NH(C₁-C₄)alkyl or -(CH₂)_n-C(=O)N(C₁-C₄)alkyl (C₁-C₄ alkyl), CN, NO₂, -OR¹⁶;

R³ and R⁵ are independently H, alkyl C₁-C₆, substituted alkyl C₁-C₆, cycloalkyl C₁-C₆ and substituted cycloalkyl C₁-C₆, (C₂-C₄)alkyl-O-(C₁-C₄)alkyl, (C₂-C₄)alkyl-NH(C₁-C₄)alkyl, (C₂-C₄)alkyl-N(C₁-C₄)alkyl(C₁-C₄ alkyl), (C₁-C₄)alkyl-heterocyclic;

R⁶ and R⁷ are independently C₁-C₄ alkyl;

each R¹³, R¹⁴, and R¹⁵ are independently selected from H, -C₁-C₄ alkyl, -(C₂-C₄)alkyl-O-(C₁-C₄)alkyl, -(CH₂)_v-NR¹⁶R¹⁷, or a 4- to 7-membered heterocyclic group; or R¹³ and R¹⁴ when in -NR¹³R¹⁴, may optionally be connected to form a 4 to 6 membered heterocyclic group, which heterocyclic group optionally comprises from 1 to 3 further hetero moieties selected from N, S, O and -C(=O);

R¹⁶ and R¹⁷ are independently H, C₁-C₆ alkyl or together may form a 4- to 7-membered heterocyclic group;

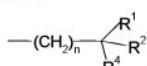
k is an integer selected from zero, 1, 2, 3, 4, and 5; and

v is an integer selected from 2, 3, 4, and 5; and

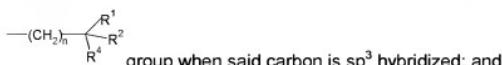
n is an integer selected from zero, 1, 2, 3, 4, and 5;

and pharmaceutically acceptable salts thereof;

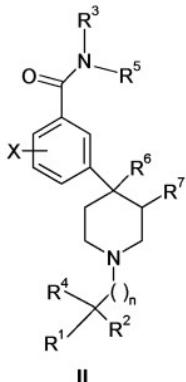
with the proviso that;



a) in said group, when n is 0, R¹, R² or R⁴ cannot be a heteroatom or contain a heteroatom which is directly linked to the carbon of said



- b) R¹³ and R¹⁴ cannot be H in a -NHS(=O)₂R¹⁴ or a -SO₂R¹³ group.
2. (Original) The compound according to claim 1 represented by the chemical structure II:



Wherein each of X, R¹, R², R³, R⁴, R⁵, R⁶, R⁷ and n is represented as described in claim 1 and the preferred relative stereochemistry between R⁶ and R⁷ is *trans*.

3. (Original) The compound according to claim 2 wherein R³ and R⁵ are H.
4. (Original) The compound according to claim 2 wherein X is H.
5. (Original) The compound according to claim 2 wherein R⁶ and R⁷ are each CH₃.
6. (Original) The compound according to claim 2 wherein n is 1, 2 or 3.
7. (Original) The compound according to claim 2 wherein R⁴ is OH, CH₂OH, NH₂, NHCOCH₃ or CN.
8. (Cancelled).
9. (Original) The compound according to claim 6 wherein n is 1.
10. (Original) The compound according to claim 7 wherein R⁴ is OH.
- 11-19. (Cancelled).
20. (Currently Amended) A pharmaceutical composition comprising an effective amount of a compound according to any of claims 1-7 and 9-10 1-19 in combination with a pharmaceutically acceptable carrier, excipient or additive.
21. (Currently Amended) A method of treating in a mammal, in need thereof, a disease state, disorder or condition mediated by an opioid receptor or receptors which

method comprises administering to said mammal an amount of a compound according to any of claims 1-7 and 9-10 1-19 effective in modulating an opioid receptor or receptors.

22. (Currently Amended) A method of treating in a mammal, in need thereof, a disease state, disorder or condition selected from the group consisting of irritable bowel syndrome, constipation, nausea, vomiting, pruritic dermatoses, psoriasis; eczema; an insect bite; an eating disorder, depression, anxiety, schizophrenia; drug addiction, an opioid overdose, sexual dysfunction, stroke, head trauma, traumatic brain injury, spinal damage, Parkinson's disease, Alzheimer's disease, age-related cognitive decline and Attention Deficit and Hyperactivity Disorder which method comprises administering to said mammal an amount of a compound according to any of claims 1-7 and 9-10 1-19 effective in treating said disease state, disorder or condition.

23. (Currently Amended) A method of treating in a mammal, in need thereof, a disease state, disorder or condition selected from the group consisting of irritable bowel syndrome, drug addiction, depression, anxiety, schizophrenia and eating disorders which method comprises administering to said mammal an amount of a compound according to any of claims 1-7 and 9-10 1-19 effective in treating said disease state, disorder or condition.

24. (Currently amended) A method of treating in a mammal, in need thereof, a disease state, disorder or condition selected from the group consisting of allergic dermatitis, contact dermatitis, anorexia, bulimia, obesity, alcohol addiction, amphetamine addiction, cocaine addiction, morphine addiction, opium addiction, heroin addiction, erectile dysfunction and impotence, which method comprises administering to said mammal an effective amount of a compound according to any of claims 1-7 and 9-10 1-19 for treating said disease state, disorder or condition.

25. – 27. (Cancelled)

28. (Original) A compound according to claim 1 wherein one or more atoms thereof have an atomic mass or mass number different from the atomic mass or mass number usually found in nature, or a pharmaceutically acceptable salt of such compound.

29. (Cancelled).